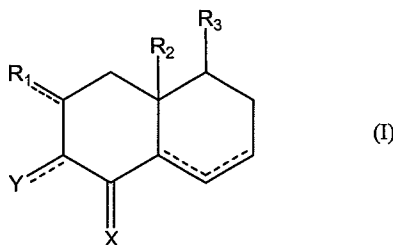


## AMENDMENTS TO THE CLAIMS

### 1.-25. (Canceled)

26. (Original) A method for controlling pests, said method comprising exposing said pests to a pest-controlling effective amount of a compound of formula (I) or a tautomer thereof or a composition comprising at least one compound of formula (I) or a tautomer thereof:



wherein:

X is selected from O, S or N-R<sub>4</sub>;

when ----- is a single bond attached to Y, Y is selected from the group consisting of H, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>halo, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>OR<sub>5</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>SR<sub>5</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=O)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=S)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>N(R<sub>4</sub>)<sub>2</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=NR<sub>4</sub>)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>NO<sub>2</sub> and [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>NR<sub>4</sub>OR<sub>8</sub>;

when ----- is a double bond attached to Y, Y is O;

when ----- is a single bond attached to R<sub>1</sub>, R<sub>1</sub> is selected from the group consisting of H, OH, SH, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>2</sub>-C<sub>10</sub> alkynyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>8</sub>-C<sub>13</sub> arylalkenyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkenyl, C<sub>4</sub>-C<sub>10</sub> cycloalkylalkyl, C<sub>4</sub>-C<sub>10</sub> cycloalkenylalkyl, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>4</sub>-C<sub>12</sub> heterocyclylalkyl, C<sub>5</sub>-C<sub>13</sub> heterocyclylalkenyl, C<sub>1</sub>-C<sub>10</sub> alkoxy, C<sub>2</sub>-C<sub>10</sub> alkenyloxy, C<sub>1</sub>-C<sub>10</sub> alkylthio, C<sub>2</sub>-C<sub>10</sub> alkenylthio, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>halo, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=O)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=S)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>N(R<sub>4</sub>)<sub>2</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=NR<sub>4</sub>)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>NO<sub>2</sub> and [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>NR<sub>4</sub>OR<sub>8</sub>;

when ----- is a double bond attached to R<sub>1</sub>, R<sub>1</sub> is CR<sub>1a</sub>R<sub>1b</sub> wherein R<sub>1a</sub> and R<sub>1b</sub> are independently selected from C<sub>1</sub>-C<sub>10</sub>alkyl;

R<sub>2</sub> and R<sub>3</sub> are independently selected from the group consisting of H, OH, SH, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>2</sub>-C<sub>10</sub> alkynyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>8</sub>-C<sub>13</sub> arylalkenyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkenyl, C<sub>4</sub>-C<sub>10</sub> cycloalkylalkyl, C<sub>4</sub>-C<sub>10</sub> cycloalkenylalkyl, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>4</sub>-C<sub>12</sub> heterocyclylalkyl, C<sub>5</sub>-C<sub>13</sub> heterocyclylalkenyl, C<sub>1</sub>-C<sub>10</sub> alkoxy, C<sub>2</sub>-C<sub>10</sub> alkenyloxy, C<sub>1</sub>-C<sub>10</sub>

alkylthio, C<sub>2</sub>-C<sub>10</sub> alkenylthio, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>halo, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=O)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=S)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>N(R<sub>4</sub>)<sub>2</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=NR<sub>4</sub>)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>NO<sub>2</sub> and [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>NR<sub>4</sub>OR<sub>8</sub>;

each R<sub>4</sub> is independently selected from the group consisting of H, OH, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>8</sub>-C<sub>13</sub> arylalkenyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkenyl, C<sub>4</sub>-C<sub>10</sub> cycloalkylalkyl, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>4</sub>-C<sub>12</sub> heterocyclalkyl, C<sub>5</sub>-C<sub>13</sub> heterocyclalkenyl, C<sub>1</sub>-C<sub>10</sub> alkoxy and C<sub>2</sub>-C<sub>10</sub> alkenyloxy;

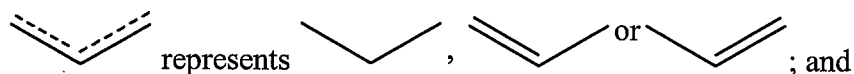
R<sub>5</sub> is selected from the group consisting of H, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>8</sub>-C<sub>13</sub> arylalkenyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkenyl, C<sub>4</sub>-C<sub>10</sub> cycloalkylalkyl, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>4</sub>-C<sub>12</sub> heterocyclalkyl, C<sub>5</sub>-C<sub>13</sub> heterocyclalkenyl, (C=O)R<sub>6</sub>, PO<sub>3</sub>R<sub>8</sub>, SO<sub>3</sub>R<sub>8</sub> and SO<sub>2</sub>R<sub>8</sub>;

R<sub>6</sub> is selected from the group consisting of H, OH, C<sub>1</sub>-C<sub>10</sub> alkoxy, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyloxy, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>6</sub>-C<sub>10</sub> aryloxy, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkenyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyloxy, C<sub>3</sub>-C<sub>6</sub> cycloalkenyloxy, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>3</sub>-C<sub>10</sub> heterocyclloxy, C<sub>1</sub>-C<sub>10</sub> alkylthio, C<sub>1</sub>-C<sub>10</sub> alkenylthio, C<sub>6</sub>-C<sub>10</sub> arylthio, C<sub>3</sub>-C<sub>6</sub> cycloalkylthio, and C<sub>3</sub>-C<sub>10</sub> heterocyclthio;

R<sub>7</sub> is selected from the group consisting of H, halogen, OR<sub>5</sub>, SR<sub>5</sub>, N(R<sub>4</sub>)<sub>2</sub>, (C=O)R<sub>6</sub>, (C=S)R<sub>6</sub>, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>4</sub>-C<sub>12</sub> heterocyclalkyl, C<sub>4</sub>-C<sub>10</sub> cycloalkylalkyl, C<sub>8</sub>-C<sub>13</sub> arylalkenyl, C<sub>5</sub>-C<sub>13</sub> heterocyclalkenyl, and NO<sub>2</sub>;

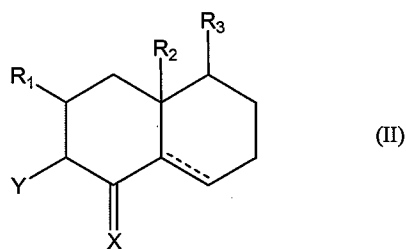
R<sub>8</sub> is selected from the group consisting of H, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>8</sub>-C<sub>13</sub> arylalkenyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkenyl, C<sub>4</sub>-C<sub>10</sub> cycloalkylalkyl, C<sub>5</sub>-C<sub>10</sub> cycloalkylalkenyl, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>4</sub>-C<sub>12</sub> heterocyclalkyl and C<sub>5</sub>-C<sub>13</sub> heterocyclalkenyl;

n is 0 or an integer selected from 1 to 5;



wherein each alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl and heterocyclyl group is optionally substituted.

27. (Currently amended) A method according to claim 263 wherein the compound of formula (I) is a compound of formula (II):



wherein:

X is selected from the group consisting of O, S or N-R<sub>4</sub>;

Y is selected from the group consisting of H, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>halo, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>OR<sub>5</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>SR<sub>5</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=O)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=S)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>N(R<sub>4</sub>)<sub>2</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=NR<sub>4</sub>)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>NO<sub>2</sub> and [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>NR<sub>4</sub>OR<sub>8</sub>;

R<sub>1</sub>, R<sub>2</sub> and R<sub>3</sub> are independently selected from the group consisting of H, OH, SH, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>2</sub>-C<sub>10</sub> alkynyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>8</sub>-C<sub>13</sub> arylalkenyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkenyl, C<sub>4</sub>-C<sub>10</sub> cycloalkylalkyl, C<sub>4</sub>-C<sub>10</sub> cycloalkenylalkyl, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>4</sub>-C<sub>12</sub> heterocyclylalkyl, C<sub>5</sub>-C<sub>13</sub> heterocyclylalkenyl, C<sub>1</sub>-C<sub>10</sub> alkoxy, C<sub>2</sub>-C<sub>10</sub> alkenyloxy, C<sub>1</sub>-C<sub>10</sub> alkylthio, C<sub>2</sub>-C<sub>10</sub> alkenylthio, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>halo, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=O)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=S)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>N(R<sub>4</sub>)<sub>2</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=NR<sub>4</sub>)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>NO<sub>2</sub> and [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>NR<sub>4</sub>OR<sub>8</sub>;

each R<sub>4</sub> is independently selected from the group consisting of H, OH, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>8</sub>-C<sub>13</sub> arylalkenyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkenyl, C<sub>4</sub>-C<sub>10</sub> cycloalkylalkyl, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>4</sub>-C<sub>12</sub> heterocyclylalkyl, C<sub>5</sub>-C<sub>13</sub> heterocyclylalkenyl, C<sub>1</sub>-C<sub>10</sub> alkoxy and C<sub>2</sub>-C<sub>10</sub> alkenyloxy;

R<sub>5</sub> is selected from the group consisting of H, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>8</sub>-C<sub>13</sub> arylalkenyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkenyl, C<sub>4</sub>-C<sub>10</sub> cycloalkylalkyl, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>4</sub>-C<sub>12</sub> heterocyclylalkyl, C<sub>5</sub>-C<sub>13</sub> heterocyclylalkenyl, (C=O)R<sub>6</sub>, PO<sub>3</sub>R<sub>8</sub>, SO<sub>3</sub>R<sub>8</sub> and SO<sub>2</sub>R<sub>8</sub>;

R<sub>6</sub> is selected from the group consisting of H, OH, C<sub>1</sub>-C<sub>10</sub> alkoxy, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyloxy, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>6</sub>-C<sub>10</sub> aryloxy, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkenyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyloxy, C<sub>3</sub>-C<sub>6</sub> cycloalkenyloxy, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>3</sub>-C<sub>10</sub> heterocycliloxy, C<sub>1</sub>-C<sub>10</sub> alkylthio, C<sub>1</sub>-C<sub>10</sub> alkenylthio, C<sub>6</sub>-C<sub>10</sub> arylthio, C<sub>3</sub>-C<sub>6</sub> cycloalkylthio, and C<sub>3</sub>-C<sub>10</sub>

heterocyclylthio;

R<sub>7</sub> is selected from the group consisting of H, halogen, OR<sub>5</sub>, SR<sub>5</sub>, N(R<sub>4</sub>)<sub>2</sub>, (C=O)R<sub>6</sub>, (C=S)R<sub>6</sub>, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>4</sub>-C<sub>12</sub> heterocyclalkyl, C<sub>4</sub>-C<sub>10</sub> cycloalkylalkyl, C<sub>8</sub>-C<sub>13</sub> arylalkenyl, C<sub>5</sub>-C<sub>13</sub> heterocyclalkenyl, and NO<sub>2</sub>;

R<sub>8</sub> is selected from the group consisting of H, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>8</sub>-C<sub>13</sub> arylalkenyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkenyl, C<sub>4</sub>-C<sub>10</sub> cycloalkylalkyl, C<sub>5</sub>-C<sub>10</sub> cycloalkylalkenyl, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>4</sub>-C<sub>12</sub> heterocyclalkyl and C<sub>5</sub>-C<sub>13</sub> heterocyclalkenyl;

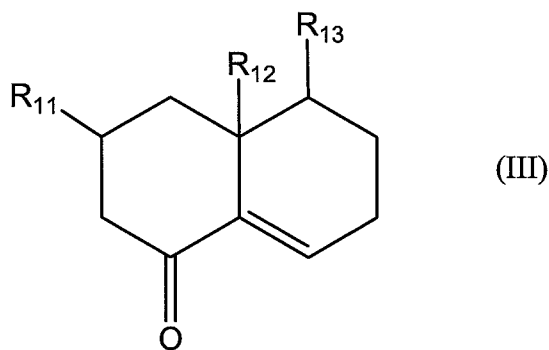
n is 0 or an integer selected from 1 to 5;

----- represents a single or double bond; and

wherein each alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl and heterocyclyl group is optionally substituted.

28. (Canceled)

29. (Currently amended) A method according to claim 263, wherein at least one compound of formula (I) is a compound of formula (III):



wherein

R<sub>11</sub> is selected from the group consisting of C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>6</sub>-C<sub>12</sub> heteroarylalkyl and C<sub>2</sub>-C<sub>10</sub> alkenyloxy wherein each C<sub>2</sub>-C<sub>10</sub> alkenyl or C<sub>2</sub>-C<sub>10</sub> alkenyloxy is optionally substituted with 1 to 3 halo, hydroxy, thiol or nitro groups; and

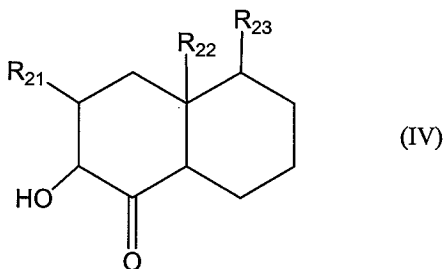
R<sub>12</sub> and R<sub>13</sub> are independently selected from the group consisting of H, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>2</sub>-C<sub>10</sub> alkynyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>3</sub>-C<sub>10</sub> cycloalkyl, C<sub>5</sub>-C<sub>10</sub> heteroaryl, C<sub>6</sub>-C<sub>12</sub> heteroarylalkyl and C<sub>1</sub>-C<sub>10</sub> alkoxy, wherein each C<sub>1</sub>-C<sub>10</sub> alkyl and C<sub>1</sub>-C<sub>10</sub> alkoxy is optionally substituted with 1 to 3 halo, hydroxy, thiol or nitro groups.

30. **(Currently amended)** A method according to claim 295, wherein R<sub>11</sub> is C<sub>2</sub>-C<sub>10</sub> alkenyl optionally substituted with a hydroxy, nitro or thiol group or 1 to 3 halo groups, and R<sub>12</sub> and R<sub>13</sub> are independently selected from C<sub>1</sub>-C<sub>10</sub> alkyl optionally substituted with a hydroxy, nitro or thiol group or 1 to 3 halo groups.

31. **(Currently amended)** A method according to claim 263 wherein at least one compound of formula (I) is eremophilone.

32. **(Canceled)**

33. **(Currently amended- Withdrawn)** A method according to claim 263 wherein at least one compound of formula (I) is a compound of formula (IV):



wherein R<sub>21</sub>, R<sub>22</sub> and R<sub>23</sub> are independently selected from the group consisting of H, OH, SH, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>2</sub>-C<sub>10</sub> alkynyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>8</sub>-C<sub>13</sub> arylalkenyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkenyl, C<sub>4</sub>-C<sub>10</sub> cycloalkylalkyl, C<sub>4</sub>-C<sub>10</sub> cycloalkenylalkyl, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>4</sub>-C<sub>12</sub> heterocyclylalkyl, C<sub>5</sub>-C<sub>13</sub> heterocyclylalkenyl, C<sub>1</sub>-C<sub>10</sub> alkoxy, C<sub>2</sub>-C<sub>10</sub> alkenyloxy, C<sub>1</sub>-C<sub>10</sub> alkylthio, C<sub>2</sub>-C<sub>10</sub> alkenylthio, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>halo, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=O)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=S)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>N(R<sub>4</sub>)<sub>2</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>(C=NR<sub>4</sub>)R<sub>6</sub>, [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>NO<sub>2</sub> and [C(R<sub>7</sub>)<sub>2</sub>]<sub>n</sub>NR<sub>4</sub>OR<sub>8</sub>;

each R<sub>4</sub> is independently selected from the group consisting of H, OH, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>8</sub>-C<sub>13</sub> arylalkenyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkenyl, C<sub>4</sub>-C<sub>10</sub> cycloalkylalkyl, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>4</sub>-C<sub>12</sub> heterocyclylalkyl, C<sub>5</sub>-C<sub>13</sub> heterocyclylalkenyl, C<sub>1</sub>-C<sub>10</sub> alkoxy and C<sub>2</sub>-C<sub>10</sub> alkenyloxy;

R<sub>6</sub> is selected from the group consisting of H, OH, C<sub>1</sub>-C<sub>10</sub> alkoxy, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyloxy, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>6</sub>-C<sub>10</sub> aryloxy, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkenyl,

C<sub>3</sub>-C<sub>6</sub> cycloalkyloxy, C<sub>3</sub>-C<sub>6</sub> cycloalkenyloxy, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>3</sub>-C<sub>10</sub> heterocyclyloxy, C<sub>1</sub>-C<sub>10</sub> alkylthio, C<sub>1</sub>-C<sub>10</sub> alkenylthio, C<sub>6</sub>-C<sub>10</sub> arylthio, C<sub>3</sub>-C<sub>6</sub> cycloalkylthio, and C<sub>3</sub>-C<sub>10</sub> heterocyclylthio;

R<sub>7</sub> is selected from the group consisting of H, halogen, OR<sub>5</sub>, SR<sub>5</sub>, N(R<sub>4</sub>)<sub>2</sub>, (C=O)R<sub>6</sub>, (C=S)R<sub>6</sub>, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>4</sub>-C<sub>12</sub> heterocyclylalkyl, C<sub>4</sub>-C<sub>10</sub> cycloalkylalkyl, C<sub>8</sub>-C<sub>13</sub> arylalkenyl, C<sub>5</sub>-C<sub>13</sub> heterocyclylalkenyl, and NO<sub>2</sub>;

R<sub>8</sub> is selected from the group consisting of H, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>8</sub>-C<sub>13</sub> arylalkenyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkenyl, C<sub>4</sub>-C<sub>10</sub> cycloalkylalkyl, C<sub>5</sub>-C<sub>10</sub> cycloalkylalkenyl, C<sub>3</sub>-C<sub>10</sub> heterocyclyl, C<sub>4</sub>-C<sub>12</sub> heterocyclylalkyl and C<sub>5</sub>-C<sub>13</sub> heterocyclylalkenyl; and

n is 0 or an integer selected from 1 to 5;

wherein each alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl and heterocyclyl group is optionally substituted.

34. **(Currently amended- Withdrawn)** A method according to claim 338 wherein R<sub>21</sub> is selected from the group consisting of C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>6</sub>-C<sub>12</sub> heteroarylalkyl and C<sub>2</sub>-C<sub>10</sub> alkenyloxy wherein each C<sub>2</sub>-C<sub>10</sub> alkenyl or C<sub>2</sub>-C<sub>10</sub> alkenyloxy is optionally substituted with 1 to 3 halo, hydroxy, thiol or nitro groups; and

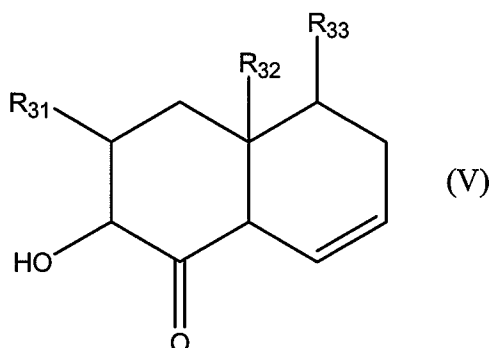
R<sub>22</sub> and R<sub>23</sub> are independently selected from the group consisting of H, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>2</sub>-C<sub>10</sub> alkynyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>3</sub>-C<sub>10</sub> cycloalkyl, C<sub>5</sub>-C<sub>10</sub> heteroaryl, C<sub>6</sub>-C<sub>12</sub> heteroarylalkyl and C<sub>1</sub>-C<sub>10</sub> alkoxy, wherein each C<sub>1</sub>-C<sub>10</sub> alkyl and C<sub>1</sub>-C<sub>10</sub> alkoxy is optionally substituted with 1 to 3 halo, hydroxy, thiol or nitro groups.

35. **(Currently amended- Withdrawn)** A method according to claim 349 wherein R<sub>21</sub> is C<sub>2</sub>-C<sub>10</sub> alkenyl, optionally substituted with a hydroxy, thiol or nitro group or 1 to 3 halo groups, and R<sub>22</sub> and R<sub>23</sub> are independently selected from C<sub>1</sub>-C<sub>10</sub> alkyl, optionally substituted with a hydroxy, thiol or nitro group or 1 to 3 halo groups.

36. **(Currently amended- Withdrawn)** A method according to claim 263 wherein at least one compound of formula (I) is 8-hydroxy-1(10) dihydroeremophilone.

37. **(Canceled)**

38. **(Currently amended- Withdrawn)** A method according to claim 263 comprising at least one compound of formula (V):



wherein R<sub>31</sub> is selected from the group consisting of C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>6</sub>-C<sub>12</sub> heteroarylalkyl and C<sub>2</sub>-C<sub>10</sub> alkenyloxy wherein each C<sub>2</sub>-C<sub>10</sub> alkenyl or C<sub>2</sub>-C<sub>10</sub> alkenyloxy is optionally substituted with 1 to 3 halo, hydroxy, thiol or nitro groups; and

R<sub>32</sub> and R<sub>33</sub> are independently selected from the group consisting of H, C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>2</sub>-C<sub>10</sub> alkenyl, C<sub>2</sub>-C<sub>10</sub> alkynyl, C<sub>6</sub>-C<sub>10</sub> aryl, C<sub>7</sub>-C<sub>12</sub> arylalkyl, C<sub>3</sub>-C<sub>10</sub> cycloalkyl, C<sub>5</sub>-C<sub>10</sub> heteroaryl, C<sub>6</sub>-C<sub>12</sub> heteroarylalkyl and C<sub>1</sub>-C<sub>10</sub> alkoxy, wherein each C<sub>1</sub>-C<sub>10</sub> alkyl and C<sub>1</sub>-C<sub>10</sub> alkoxy is optionally substituted with 1 to 3 halo, hydroxy, thiol or nitro groups.

39. **(Currently amended- Withdrawn)** A method according to claim 3812 wherein R<sub>31</sub> is C<sub>2</sub>-C<sub>10</sub> alkenyl optionally substituted with a hydroxy, nitro or thiol group or 1 to 3 halo groups, and R<sub>32</sub> and R<sub>33</sub> are independently selected from C<sub>1</sub>-C<sub>10</sub> alkyl optionally substituted with a hydroxy, nitro or thiol group or 1 to 3 halo groups.

40. **(Currently amended- Withdrawn)** A method according to claim 263 wherein at least one compound of formula (I) is 8-hydroxyeremophila-1,11-dienone.

41. **(Currently amended)** A method according to claim 263 wherein the composition comprises an extract containing at least one compound of formula (I) obtained from a volatile oil bearing plant from the Myoporaceae family.

42. **(Canceled)**

43. **(Canceled)**

44. **(Currently amended)** A method according to claim 263 wherein the pest-controlling effective amount is a pesticidally effective amount.

45. **(Currently amended)** A method according to claim 263 wherein the pest-controlling effective amount is a pest-repelling effective amount.

46. **(Currently amended)** A method according to claim 263 wherein the pest-controlling effective amount is a antifeedant effective amount.
47. **(Currently amended)** A method according to claim 263 wherein the pests are selected from the group consisting of insects, arachnids, helminths and molluscs.
48. **(Currently amended)** A method according to claim 263 wherein the pests are selected from the group consisting of termites, earwigs, cockroaches and wood borer beetles and their larvae.
49. **(Currently amended)** A method according to claim 263 wherein the pests are wood associated pests.
50. **(Currently amended)** A method according to claim 4924 wherein the wood associated pests are selected from the group consisting of termites and wood borer beetles.
51. **(Currently amended)** A method according to claim 5022 wherein the wood associated pests are termites.
52. **(Currently amended)** A method according to claim 263 wherein pests are exposed to the pest-controlling effective amount of a compound of formula (I) or a composition comprising at least one compound of formula (I) by applying the compound or composition to a site of infestation, a potential site of infestation, a habitat of the pest or a potential habitat of the pest.
53. **(Currently amended)** A method according to claim 5224 wherein the compound or composition is applied to a surface or impregnated into a material or article of manufacture.
54. **(Currently amended)** A method according to claim 5325 wherein the compound or composition is applied to a surface by spraying, coating or painting the surface.
55. **(Currently amended)** A method according to claim 5426 wherein the surface is a soil surface, timber, buildings, wooden articles of manufacture or a physical barrier.
56. **(Currently amended)** A method according to claim 5527 wherein the material or article of manufacture is soil, timber, timber or wooden products or buildings or parts of buildings.
57. **(Currently amended)** A method according to claim 5224 wherein the compound or composition is applied in a band or furrow around a site of infestation or potential infestation or is mixed with a layer of soil at a site of infestation or a potential site of infestation.



58.-78. (Canceled)

79. (Currently amended) A method of combating an already existing wood associated pest infestation comprising applying at least one compound of formula (I) or a tautomer thereof or a composition comprising at least one compound of formula (I) or a tautomer thereof ~~a composition according to claim 1~~ to a wood associated pest affected surface, wherein the compound of formula (I) is as defined in Claim 26.

80.-82. (Canceled)